Translation

PATENT COOPERATION TREATY

PCT/MX2003/000093

PCT

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference SOPH.PCT/003	FOR FURTHER ACTION See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)			
International application No. PCT/MX2003/000093	International filing date (da		Priority date (day/month/year)	
PCT/MX2003/000093 30 October 2003 (30.10.2003) 13 December 2002 (13.12.2002) International Patent Classification (IPC) or national classification and IPC A61K 31/7004, 9/08			13 December 2002 (13.12.2002)	
Applicant ЛМЕNEZ BAYARDO, Arturo				
 This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36. 				
2. This REPORT consists of a total of	5 sheets, inclu	ding this cover s	sheet.	
This report is also accompanied by ANNEXES, i.e., sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).				
These annexes consist of a to	tal of 3 sheets			
3. This report contains indications rela	ting to the following items:			
I Basis of the report				
II Priority	II Priority			
III Non-establishment o	of opinion with regard to nove	lty, inventive st	ep and industrial applicability	
IV Lack of unity of inve	ention			
V Reasoned statement citations and explana	Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement			
VI Certain documents of	ited			
VII Certain defects in th	e international application			
VIII Certain observations on the international application				
Date of submission of the demand Date of completion of this report		of this report		
05 October 2004 (05.10.2004)		17 Fe	ebruary 2005 (17.02.2005)	
Name and mailing address of the IPEA/ES		orized officer		
Facsimile No.		Telephone No.		

Form PCT/IPEA/409 (cover sheet) (July 1998)

International application No.

PCT/MX2003/000093

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1.	With	regard to	to the elements of the international application:*	· · · · · · · · · · · · · · · · · · ·
		the inte	ternational application as originally filed	
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2.	the in	e elemen	to the language, all the elements marked above were available or furnished to this Authority in the conal application was filed, unless otherwise indicated under this item. Into were available or furnished to this Authority in the following language	
	H		nguage of a translation furnished for the purposes of international search (under Rule 23.1(b)).	
	H		nguage of publication of the international application (under Rule 48.3(b)).	
		or 55.3		
3.	With	regard minary ex	I to any nucleotide and/or amino acid sequence disclosed in the international application examination was carried out on the basis of the sequence listing:	, the international
			ined in the international application in written form.	
	Щ	filed to	ogether with the international application in computer readable form.	
		furnish	hed subsequently to this Authority in written form.	
		furnish	hed subsequently to this Authority in computer readable form.	
		The sta	statement that the subsequently furnished written sequence listing does not go beyond the ational application as filed has been furnished.	disclosure in the
		The sta	tatement that the information recorded in computer readable form is identical to the written sefurnished.	equence listing has
4.		The am	mendments have resulted in the cancellation of:	
			the description, pages	
		<u> </u>	the claims, Nos.	
			the drawings, sheets/fig	
5.		This rep	eport has been established as if (some of) the amendments had not been made, since they have been it the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)).**	en considered to go
-	Replacin this	s report	sheets which have been furnished to the receiving Office in response to an invitation under Article t as "originally filed" and are not annexed to this report since they do not contain amenda	: 14 are referred to ments (Rule 70.16
		•	nent sheet containing such amendments must be referred to under item 1 and annexed to this report.	

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Supplemental Box

(To be used when the space in any of the preceding boxes is not sufficient)

Continuation of: III.1

Claims 8 to 12 relate to subject matter which, in the opinion of this Authority, is subject to the provisions of PCT Rule 67.1(iv), concerning methods for the treatment of the human or animal body by surgery or therapy. Consequently no opinion on the industrial applicability of the subject matter of these claims will be established (PCT Article 34(4)(a)).

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V .	Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability
	citations and explanations supporting such statement

1.	Statement			
	Novelty (N)	Claims	1-17	YES
		Claims		NO
	Inventive step (IS)	Claims	1-17	YES
		Claims	···-	NO NO
	Industrial applicability (IA)	Claims	1-7, 13-17	YES
		Claims		NO

2. Citations and explanations (Rule 70.7)

Document	Publication or	Publication date	
	identification no.		
D01	ES 2067418 A	16 March 1995	
D02	WO 9718835A	29 May 1997	
D03	WO 0051620 A	8 September 2000	
D04	EP 868909 A	07 October 1998	

The present invention concerns an intravitreally injectable solution for treating vitreous haemorrhages which comprises an active ingredient, mannitol, and a carrier solution (Sophisen), which comprises polyoxylic stearate, dehydrated disodium edate, sodium chloride, boric acid, sorbic acid, sodium bisulphite and distilled water. The invention also concerns the method for preparing this solution.

Document D1 concerns the use of human recombinant interleukin-1 beta in the production of medicaments for treating human intravitreal haemorrhages in the form of intravitreal injections which are intended to be administered transconjunctivally.

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Document D2 concerns the use of enzymes for eliminating vitreal haemorrhaging. These enzymes are in the form of a liquid solution which is injected into the vitreous humour.

Document D3 concerns methods for treating ocular disorders, including the acceleration of the elimination of blood from the vitreous humour. Urea, urea derivatives, non-steroid anti-inflammatories, etc. are applied by means of intravitreal injection.

Document D4 uses the same carrier means as the application, together with various therapeutic agents for topical ophthalmic application.

Therefore, none of the citations discloses mannitol as active ingredient together with the carrier solution whose composition is described in claim 1. The subject matter of claims 1 to 17 is thus considered to meet the novelty and inventive step requirements (PCT Article 33(2) and (3)).

Finally, the subject matter of claims 1 to 7 and 13 to 17 is considered to meet the industrial applicability requirements as defined in PCT Article 33(4). The PCT Contracting States have no uniform criteria as concerns the industrial applicability of claims 8 to 12. Patentability may also depend on the wording of the claims.